

# SCORE Search Results Details for Application 10578781 and Search Result 20081104\_154454\_us-10-578-781-1.rng.

<a href="#">Score Home</a>	<a href="#">Retrieve Application</a>	<a href="#">SCORE System</a>	<a href="#">SCORE</a>	<a href="#">Comments /</a>
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This page gives you Search Results detail for the Application 10578781 and Search Result 20081104\_154454\_us-10-578-781-1.rng.

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GenCore version 6.3  
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QM nucleic - nucleic search, using sw model

Run on: November 4, 2008, 17:10:32 ; Search time 243 Seconds  
(without alignments)  
44258.760 Million cell updates/sec

Title: US-10-578-781-1  
Perfect score: 756  
Sequence: 1 at gggg gccgat at caaaaa.....aggagcaaat t gaacat t ag 756

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 11806651 seqs, 7113014948 residues

Total number of hits satisfying chosen parameters: 23613302

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : N\_Geneseq\_200808: \*

- 1: geneseqn1980s: \*
- 2: geneseqn1990s: \*
- 3: geneseqn2000: \*
- 4: geneseqn2001a: \*
- 5: geneseqn2001b: \*
- 6: geneseqn2002a: \*
- 7: geneseqn2002b: \*
- 8: geneseqn2003a: \*
- 9: geneseqn2003b: \*
- 10: geneseqn2003c: \*
- 11: geneseqn2003d: \*
- 12: geneseqn2004a: \*
- 13: geneseqn2004b: \*
- 14: geneseqn2004c: \*
- 15: geneseqn2004d: \*
- 16: geneseqn2004e: \*
- 17: geneseqn2004f: \*
- 18: geneseqn2005a: \*
- 19: geneseqn2005b: \*
- 20: geneseqn2005c: \*
- 21: geneseqn2006a: \*

22: geneseqn2006b: \*  
 23: geneseqn2006c: \*  
 24: geneseqn2006d: \*  
 25: geneseqn2007a: \*  
 26: geneseqn2007b: \*  
 27: geneseqn2007c: \*  
 28: geneseqn2007d: \*  
 29: geneseqn2008: \*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARY

Result No.	Score	% Query Match	Length	DB	ID	Description
1	756	100.0	756	18	AEA00728	Aea00728 Brevibacillus
2	273.4	36.2	665	6	ABK78469	Abk78469 Bacillus
3	192.2	25.4	1438	12	ADN60500	Adn60500 B. lichen
4	178.2	23.6	777	21	Aeh93992	Aeh93992 StgG codi
5	173.8	23.0	381	6	ABK74144	Abk74144 Bacillus
6	147	19.4	631	6	ABK74048	Abk74048 Bacillus
7	78	10.3	1110	12	ADH97013	Adh97013 S. pneumo
8	78	10.3	1110	18	AEC13368	Aec13368 Streptococ
9	76.4	10.1	915	12	ADK44581	Adk44581 Streptococ
10	76.4	10.1	915	21	AEJ68509	Aej68509 Streptococ
11	76.4	10.1	915	21	AEJ75484	Aej75484 Streptococ
12	76.4	10.1	915	21	AEJ82844	Aej82844 Streptococ
13	76.4	10.1	915	21	AEL05163	Ael05163 Streptococ
14	76.4	10.1	915	21	AEL12413	Ael12413 Streptococ
15	76.4	10.1	915	21	AEL50821	Ael50821 Streptococ
16	76.4	10.1	915	25	AEM07844	Aem07844 Streptococ
17	76.4	10.1	915	25	AEM66066	Aem66066 Streptococ
18	76.4	10.1	915	25	AEM66645	Aem66645 Streptococ
19	76.4	10.1	915	25	AGI20663	Agj20663 Streptococ
20	76.4	10.1	915	25	AEN48202	Aen48202 Streptococ
21	76.4	10.1	915	25	AEN55537	Aen55537 Streptococ
22	76.4	10.1	915	25	AEN40434	Aen40434 Streptococ
23	76.4	10.1	915	25	AGI76906	Agj76906 Streptococ
24	76.4	10.1	915	25	AEN08741	Aen08741 Streptococ
25	76.4	10.1	915	25	AGV09876	Agv09876 Streptococ
26	76.4	10.1	915	25	AGV21123	Agv21123 Streptococ
27	76.4	10.1	915	25	AJE78366	Aej78366 Streptococ
28	76.4	10.1	915	25	AJE70154	Aej70154 Streptococ
29	76.4	10.1	915	25	AJE86340	Aej86340 Streptococ
30	76.4	10.1	915	25	AJE95472	Aej95472 Streptococ
31	76.4	10.1	915	25	AJE51089	Aej51089 Streptococ
32	76.4	10.1	915	25	AGV40930	Agv40930 Streptococ
33	76.4	10.1	915	25	AGV46355	Agv46355 Streptococ
34	76.4	10.1	915	25	AJF01903	Ajf01903 Streptococ
35	76.4	10.1	915	25	AJF07993	Ajf07993 Streptococ
36	76.4	10.1	915	25	AJF13317	Ajf13317 Streptococ
37	76.4	10.1	915	25	AJF53284	Ajf53284 Streptococ
38	76.4	10.1	915	25	AJF18960	Ajf18960 Streptococ
39	76.4	10.1	915	25	AJG97788	Ajf97788 Streptococ
40	76.4	10.1	915	25	ALK14103	Alk14103 Streptococ
41	76.4	10.1	915	25	ALT08207	Alt08207 Streptococ
42	76.4	10.1	915	25	ANK69294	Ank69294 Streptococ
43	76.4	10.1	915	25	ANJ76477	Anj76477 Streptococ
44	76.4	10.1	915	25	ANK74634	Ank74634 Streptococ
45	76.4	10.1	915	26	ANN03548	Ann03548 Streptococ

## ALIGNMENTS

RESULT 1

AEA00728

ID AEA00728 standard; DNA; 756 BP.

XX

AC

XX

DT

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DE

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KW

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CS

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FT

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XX

SQ

28-JUL-2005 (first entry)

Brevibacillus choshinensis DNA #1.

Cell culture; Brevibacillus choshinensis; gene; ds.

Brevibacillus choshinensis.

Key Location/Qualifiers

CDS 1..756

/\*tag= a

/product= "B. choshinensis protein #1"

W02005045005-A1.

19-MAY-2005.

08-NOV-2004; 2004W0-JP016912.

11-NOV-2003; 2003JP-00381606.

(HGET) HIGETA SHOYU KK.

Hanagata H, Nishijyo T;

WPI; 2005-366840/37.

P-PSDB; AEA00728.

New Brevibacillus choshinensis, that does not form spores and which shows low extracellular or intracellular protease activity, useful as host for producing recombinant protein.

Claim 4; SEQ ID NO 1; 103pp; Japanese.

The invention relates to a Brevibacillus choshinensis HPD31-SP3 (FERMBP-08479), which does not form spores and which has mycological characteristics such as cell size and rod shape and physiological characteristics such as negative for nitrate reduction and positive for citric acid utilization, oxidase and catalase, and showing low extracellular protease activity. The invention also relates to a transformed B. choshinensis using a vector containing the gene encoding the protein of the invention. B. choshinensis is useful as a host for producing a recombinant protein and for producing a protein by culturing a transformed host. It has decreased extracellular protein degradation activity when compared with other strains. This sequence represents B. choshinensis DNA of the invention.

Sequence 756 BP; 207 A; 166 C; 211 G; 172 T; 0 U; 0 Q her;

Query Match 100.0% Score 756; DB 18; Length 756;  
 Best Local Similarity 100.0% Pred. No. 3e-234;  
 Matches 756; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ATGGGTGGCGATATCAAAATGGAGTCAACATTTCTGAACATGAACAGTGAAGAT 60  
 Db 1 ATGGGTGGCGATATCAAAATGGAGTCAACATTTCTGAACATGAACAGTGAAGAT 60

Qy 61 TTGATAGCCAAAGACCAAGCTGGGATACGGATGACGTGAGCTTCTGGTGAATAGCAAT 120  
 Db 61 TTGATAGCCAAAGACCAAGCTGGGATACGGATGACGTGAGCTTCTGGTGAATAGCAAT 120

Qy 121 ATCAGACTGGTCTGGTCCGTGGTCCAGGGCTTTATCAACGGGGGATGAAGGGGATGAT 180  
 Db 121 ATCAGACTGGTCTGGTCCGTGGTCCAGGGCTTTATCAACGGGGGATGAAGGGGATGAT 180

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Qy      181 TTGTTTCAGATCGGTTGCATTGGCTTGCTCAAGGCGGTTGACAAGTTGGATCTTTGGTAC 240
Db      181 TTGTTTCAGATCGGTTGCATTGGCTTGCTCAAGGCGGTTGACAAGTTGGATCTTTGGTAC 240

Qy      241 GATGTGAGATTTTGGAGCTATGGGTTGCCAATGATCATGCGAGAAATTCAAGGCTTTTGTG 300
Db      241 GATGTGAGATTTTGGAGCTATGGGTTGCCAATGATCATGCGAGAAATTCAAGGCTTTTGTG 300

Qy      301 CGGATGACGGTACGGTAAAGGTGAGTGGATCGTTAAAAAGAAACAGGAAATAGGTGGG 360
Db      301 CGGATGACGGTACGGTAAAGGTGAGTGGATCGTTAAAAAGAAACAGGAAATAGGTGGG 360

Qy      361 CGATCAAGGATGAATTGTACAAGCAATTGGGCGGTGCGGCGCAAGATGCGAGAAGTGGA 420
Db      361 CGATCAAGGATGAATTGTACAAGCAATTGGGCGGTGCGGCGCAAGATGCGAGAAGTGGA 420

Qy      421 GAAGCAGTGGGAATCAAGCGGAGGAAAGTGTCTTTGGCGAAGAGGCAAGCAGAGCGCT 480
Db      421 GAAGCAGTGGGAATCAAGCGGAGGAAAGTGTCTTTGGCGAAGAGGCAAGCAGAGCGCT 480

Qy      481 TCTCCATCCATGAGACGGTTTTTGAAGATGAAGCGGATCCATCACACTGATCGATCAG 540
Db      481 TCTCCATCCATGAGACGGTTTTTGAAGATGAAGCGGATCCATCACACTGATCGATCAG 540

Qy      541 ATAGCGGATGAAGGTGTGAACAAGTGGTTTGAAGAAATTTGGCTTGAAGGACGCGATCAGC 600
Db      541 ATAGCGGATGAAGGTGTGAACAAGTGGTTTGAAGAAATTTGGCTTGAAGGACGCGATCAGC 600

Qy      601 AGGCTGAGGAGCGGTGAGCAGCTCATGCTCACTGGGCTATTACAAGGATCAGACACAG 660
Db      601 AGGCTGAGGAGCGGTGAGCAGCTCATGCTCACTGGGCTATTACAAGGATCAGACACAG 660

Qy      661 TCTGAGGTAGCAGAGCGGTCTAGGGAATTTGCGAGGTGAGGCTCTGCGCTCTGGAAGAGGT 720
Db      661 TCTGAGGTAGCAGAGCGGTCTAGGGAATTTGCGAGGTGAGGCTCTGCGCTCTGGAAGAGGT 720

Qy      721 ATCTGCTAAGGATCAAGGAGCAAAATTGAACATTAG 756
Db      721 ATCTGCTAAGGATCAAGGAGCAAAATTGAACATTAG 756

```

## RESULT 2

ABK78469

ID ABK78469 standard; DNA; 665 BP.

XX

AC ABK78469;

XX

DT 13- AUG-2002 (first entry)

XX

DE Bacillus clausii genomic sequence tag (GST) #1312.

XX

KW Differential gene expression; genomic sequenced tag; GST;

KW altered culture condition; environmental stress;

KW physiological provocation; ds.

XX

OS Bacillus clausii.

XX

PN WQ200229113-A2.

XX

PD 11- APR-2002.

XX

PF 05- OCT-2001; 2001WD-US031437.

XX

PR 06- OCT-2000; 2000US-00680598.

PR

27- MAR-2001; 2001US-0279526P.

XX

PA (NOVO) NOVOZYMES BIOTECH INC.

PA (NOVO) NOVOZYMES AS.

Query Match 36.2% Score 273.4; DB 6; Length 665;  
Best Local Similarity 67.4% Pred. No. 1.9e-77;  
Matches 399; Conservative 0; M smatches 192; Indels 1; Gaps 1;

Qy	37	CTGA	CCAA	TGAC	CAAG	TGAA	AGAT	TGAT	AGCC	AAGAG	GC	CAAG	CTGG	GATAC	GGAT	GCA	96									
Db	40	CTAT	CCGA	TAA	CAAG	TGAA	AGAT	TGAT	TGCA	AAAGAG	GC	AGGA	GGGG	ACAC	AGAA	GCA	99									
Qy	97	CGT	GAG	CTT	CTCG	TAA	TAG	CAAT	ATC	AGACT	GGT	CTG	GTG	CGG	TGTC	CGCG	CTTT	ATC	155							
Db	100	CGG	ATT	CGAT	TGTC	CA	CCAT	AAC	ACAG	CTG	CTG	CTG	TGAG	TGTT	GTT	CAAG	GGTTTT	TG	159							
Qy	157	AAC	CGCG	GGT	ATG	AAG	CGG	TAT	GAT	TGTT	TGAT	AGG	TGAT	GC	CTG	GCT	CAAG	GGC	216							
Db	160	AAT	CGCG	GGT	TAT	GAG	GC	AGAT	GAC	CTTT	CAAA	TGG	CTG	CAAT	TGTT	TAAT	TAG	CT	219							
Qy	217	GTT	GACA	AGT	TCG	AT	CTTT	TG	TAC	AGT	TG	GAG	ATTTT	TG	ACG	CTAT	GCGG	TGCA	TAT	276						
Db	220	GT	GC	ACAA	ATTT	GAC	TTT	TC	TAC	AGG	TG	GAA	ATTT	TC	ACG	TAT	TG	TG	GGG	TAT	279					
Qy	277	AT	CGG	A	GAA	ATTC	AA	CGCT	TTTTT	TG	CGG	TAT	GAC	GGT	AAG	GT	CAG	TG	AT	336						
Db	280	ATT	GGT	GAA	TCC	AAC	GGT	TTT	CT	G	GGG	TAT	G	GC	AC	AGT	GAA	AGT	AAG	CGGG	TC	339				
Qy	337	AAA	GAA	ACAG	CGA	ATA	AGG	TG	GGG	GGAT	CAAA	GGAT	GA	TTT	GT	ACA	AGCA	ATT	TGCG	CGT	396					
Db	340	AAA	GAA	TTT	AAG	CAAT	AAAA	TG	CGCA	AAAG	CAAAA	AGAC	GA	ACT	GAC	AAAA	ACG	TG	CGCG	CGG	399					
Qy	397	G	CCCC	ACG	ATG	CG	CAAG	TG	GC	AGA	GC	AGT	G	GAAT	CA	CGC	CGG	AGG	AGT	AGT	CTTT	456				
Db	400	G	CA	CGAC	CA	TTA	TAT	G	AG	TG	CGT	G	AAC	ATT	T	AG	GGT	G	AG	CGCT	GAG	GAA	TTT	GT	ATTT	459
Qy	457	G	GC	CAAG	AG	GC	CAAG	CG	CGCT	CT	CTCAT	TCAT	G	AG	AC	CGT	TTTTT	G	AAA	AT	GAC	GC	515			

```

Db      460  CCTGGAGATGCCAACGGAGCTTGTCTCAATCATGAAACGTTTATGAAATGACGGC 519
Qy      517  GATCCCATCACACTGATCGATCAGATAGCGGATGAAGGTGTGAACAAGTGGTTTGAGAAA 576
Db      520  GATCCGATTACACTTCTAGATCAAATGGGACCACTCACAAGTCAAATGGTTTGACAAG 579
Qy      577  ATTGCCTTGAAGCAAGGCATCAGCAGGCTGAGCGAGCGTGACGAGCTCATCG 628
Db      580  ATTG-CTTTAAGAAGCGATTGGACCTTGGGAGAGGAGCGGCTAATTG 630

```

## RESULT 3

ADN60500

ID ADN60500 standard; DNA; 1438 BP.

XX

AC

ADN60500;

XX

DT

01-JUL-2004 (first entry)

XX

DE

B. licheniformis sporulation related polynucleotide, seq id 172.

XX

KW

Mutant host cell; sporulation; oxidoreductase; transferase; hydrolase; lyase; isomerase; ligase; gene; ds.

XX

OS

Bacillus licheniformis.

XX

PN

W02003087148-A2.

XX

PD

23-OCT-2003.

XX

PF

25-MAR-2003; 2003W0-DK000200.

XX

PR

10-APR-2002; 2002DK-00000533.

XX

PA

(NOVO) NOVOZYMES AS.

XX

PI

Andersen JT, Jorgensen ST, Rasmussen MD, Olsen PB, Clausen IG

XX

DR

WPI; 2004-122131/12.

XX

DR

P-PSDB; ADN60501.

XX

PT

A Bacillus licheniformis mutant host cell for producing a product of

XX

interest e.g. vitamins, antibiotics and enzymes.

XX

PS

Claim 1; SEQ ID NO 172; 319pp; English.

XX

OC

The invention relates to a Bacillus licheniformis mutant host cell derived from a parent B. licheniformis host cell. The mutant host cell is mutated in one or more genes encoding one or more polypeptides involved in sporulation. The host cell comprises one or more heterologous genes present in at least two copies, encoding one or more heterologous polypeptides. The heterologous genes are stably integrated into the genome of the cell without leaving any antibiotic resistance marker genes at the site of integration. The heterologous genes are transcribed from a heterologous promoter or from an artificial promoter, and are comprised in an operon, preferably a polycistronic operon. The heterologous polypeptide is an antimicrobial peptide, or a fusion peptide comprising a peptide part which in its native form has antimicrobial activity. The heterologous polypeptide is an enzyme, preferably a secreted enzyme. The enzyme is an enzyme of a class selected from the group of enzyme classes consisting of oxidoreductases (EC 1), transferases (EC 2), hydrolases (EC 3), lyases (EC 4), isomerases (EC 5), and ligases (EC 6). The Bacillus licheniformis is useful in a process for producing at least one product of interest, comprising cultivating a B. licheniformis mutant host cell in a suitable medium whereby the said product is produced. The process further comprises isolating or purifying the product of interest. The current sequence represents a B. licheniformis sporulation related polynucleotide.

Qy		1	ATGCGTGGCGATATCAAAATGGAGTCACCATTTCTGAACAATGACC--AAGTGAAA	57
Ds		954	ATGGATGTGGAGGTTAAAAAGAAAAOCAGAACACTCAGCTTAAGAACCATGAAGTGAAA	1013
Qy		58	GATTTGATAGCCAAAGAGGCCAAGCTGGGATAOAGATGCAOGTGAAGCTTCTOCTGAATAGC	117
Ds		1014	GAACTGATTAAAAACAGGCAGAAAGCGGATCAAAAAGCAAAGCGACTCTCTCATAGAAAA	1073
Qy		118	AATATCAGACTGGTCTGGTOGTCGTCAGCGCTTTATCAACGGCGGGTATGAAGCGGAT	177
Ds		1074	AACATGGTCCTGTGTTGGTCTGTGCTTCAGGGTTTTTGAACAGAGCTATGAGCCTGAC	1133
Qy		178	GATTTGTTTCAGATGGGTTGCATTGCTTGGCTCAAGCGCGTTGACAAGTTGATCTTTTCG	237
Ds		1134	GACCTCTTTCAATGGGCTGCATGGGCTCTTGAAGTCGGTGGACAAATTCGATCTTTC	1193
Qy		238	TACGATGTGAGATTTTGAAGCTATGGGTCGCAATGATCATCGAGAAATTCAGCGCTTT	297
Ds		1194	TATGAGGTTGGTTTTTCAGCTAGCGGTTGGATGATTAAGCGAGATTGAGCGGTT	1253
Qy		298	TTGGCGGATGAAGGTAAGGTCAGTCGATGGTAAAAAGAAACAGGAATAAGGTG	357
Ds		1254	ATCAGAGATGAAGGAAAGCTCAAAAGTAGCGGCTGGCTGAAAGAACTCGGCAACAAATC	1313
Qy		358	CGCGGATCAAAAGGATGAATGTACAAGCAATTGGGCGTGGCCCCAGATGCGAGAAATG	417
Ds		1314	CGGCGGGCGAAAGAGAGCTTCCAAGTCAAAAGCGCGGATTCGAGCGGTTCAAGAAATC	1373
Qy		418	GCAAGAGCAGTGGGAATCAAGCGGAGGAAAGTGTCTTTGGCGAAGAGGCAAGCAGAGCG	477
Ds		1374	GGCGATTATCTGAAATCAGTTCAGAAAGAGGTGTTGATGGCCAGGAAGGGTGGGCTCT	1433
Qy		478	CGTTC 482	
Ds		1434	CGCTC 1438	

RESULT 4  
AEH93992

ID AEH93992 standard: cDNA: 777 BP.

XX

AC AEH93992;

XX

DT 27-JUL-2006 (first entry)

XX

DE Si gG codi ng sequence.

XX  
1991

ss; gene; protein production; sigma factor; RNA polymerase;

al kal i ne pr ot ease; f ood.

XX  
06

CS Bacillus sp.; KSMF 9865.

XX  
EW

PH	Key	Location/ Quantities
ET	CDS	1 777

ET

```

/ tag= "a
/ product= "Si gG"

```

XX

PN JP2006136221-A.

XX

PD 01-JUN-2006.

XX

PF 10- NOV- 2004; 2004JP- 00326973.





Qy	625	ATOGTCTA	CGCTG	CGCTATT	ACAAGG	ATCAG	CACAG	CTCAG	GTAG	CAGAG	CGTCTAG	GGG	684
Db	649												708
Qy	685	ATTTG	CAGG	TCCAG	GTCTG	GGTCTG	GAAAG	CGTAT	CGTAA	CGATC	AAGG	AGCAA	744
Db	709	ATATC	CAAG	CACAA	GTGTCA	AGACTT	GAAAG	CGTCA	ATCAA	CAGAT	GAATA	AAAAAT	768
Qy	745	ATTGA	749										
Db	769	ATTCA	773										

# RESULT 5

ABK74144

ID ABK74144 standard; DNA; 381 BP.

XX

AC ABK74144;

XX

DT 13- AUG- 2002 (first entry)

XX

DE Bacillus licheniformis genomic sequence tag (GST) #1435.

XX

KW Differential gene expression; genomic sequenced tag; GST;

KW altered culture condition; environmental stress;

KW physiological provocation; ds.

XX

OS Bacillus licheniformis.

XX

PN WQ200229113- A2.

XX

PD 11- APR- 2002.

XX

PF 05- OCT- 2001; 2001WD- US031437.

XX

PR 06- OCT- 2000; 2000US- 00680598.

PR 27- MAR- 2001; 2001US- 0279526P.

XX

PA (NOVO) NOVOZYMES BIOTECH INC.

PA (NOVO) NOVOZYMES AS.

XX

PI Berka R, Clausen IG;

XX

DR WPI; 2002- 416684/ 44.

XX

PT Monitoring differential expression of several genes in first Bacillus

PT cell relative to expression of same genes in one or more second Bacillus

PT cells, by using substrate containing Bacillus genomic sequenced tag

PT array.

XX

PS Claim 4; SEQ ID NO 1435; 200pp; English.

XX

CC The invention describes a method of monitoring differential expression of

CC genes in a first Bacillus cell relative to expression of the genes in

CC other Bacillus cells, comprising hybridising labelled nucleic acid probes

CC isolated from Bacillus cells to a substrate containing array of Bacillus

CC genomic sequenced tags (GST), examining the array, and determining

CC relative gene expression by an observed hybridisation reporter signal of

CC a spot in the array. The method is useful for measuring the expression of

CC genes in a first Bacillus cell relative to expression of the same genes

CC in one or more second Bacillus cells. The method is useful for monitoring

CC global expression of several genes from a Bacillus cell, discovering new

CC genes, identifying possible functions of unknown open reading frames and

CC monitoring gene copy number variation and stability. Monitoring changes

CC in expression of genes may be used to provide a representation of the way

CC in which Bacillus cells adapt to changes in culture conditions,

CC environmental stress or other physiological provocation. Extensive follow

CC -up characterisation is unnecessary, when one spot on an array equals one



PI Berka R, Clausen IG;

XX WPI; 2002-416684/44.

DR  
XX  
PT Monitoring differential expression of several genes in first *Bacillus*  
PT cell relative to expression of same genes in one or more second *Bacillus*  
PT cells, by using substrate containing *Bacillus* genomic sequenced tag  
PT array.

XX  
PS Claim 4; SEQ ID NO 1339; 200pp; English.

XX  
CC The invention describes a method of monitoring differential expression of  
CC genes in a first *Bacillus* cell relative to expression of the genes in  
CC other *Bacillus* cells, comprising hybridising labelled nucleic acid probes  
CC isolated from *Bacillus* cells to a substrate containing array of *Bacillus*  
CC genomic sequenced tags (GST), examining the array, and determining  
CC relative gene expression by an observed hybridisation reporter signal of  
CC a spot in the array. The method is useful for measuring the expression of  
CC genes in a first *Bacillus* cell relative to expression of the same genes  
CC in one or more second *Bacillus* cells. The method is useful for monitoring  
CC global expression of several genes from a *Bacillus* cell, discovering new  
CC genes, identifying possible functions of unknown open reading frames and  
CC monitoring gene copy number variation and stability. Monitoring changes  
CC in expression of genes may be used to provide a representation of the way  
CC in which *Bacillus* cells adapt to changes in culture conditions,  
CC environmental stress or other physiological provocation. Extensive follow  
CC -up characterisation is unnecessary, when one spot on an array equals one  
CC gene or one open reading frame, since sequence information is available.  
CC This sequence represents a genomic sequence tag (GST) used in the method  
CC of the invention. Note: The sequence data for this patent did not form  
CC part of the printed specification, but was obtained in electronic format  
CC directly from WPO at [ftp.wipo.int/pub/published\\_pat\\_sequences](http://ftp.wipo.int/pub/published_pat_sequences)

XX  
SQ Sequence 631 BP; 203 A; 121 C; 161 G; 146 T; 0 U; 0 Q her;

Query Match 19.4% Score 147; DB 6; Length 631;  
Best Local Similarity 56.8% Pred. No. 2.3e-38;  
Matches 270; Conservative 0; Mismatches 205; Indels 0; Gaps 0;

Qy	77	AAGCTGGGATACGGATGCAOGTGACCTTCTGTGAATAGCAATATCAGACTGGTCTGGT	136
Db	98	ATGAAGGAGACACAACAGGGAGAGAAAAGCTTGTTAAGGGCAATTTGGGGCTTTGTCCTAA	157
Qy	137	CGGTGTCGACGGCTTTTATCAAGCGGGGTATGAAGGGATGATTGTTTCAGATCGGTT	196
Db	158	GGGTGATTCAAAAGGTTTAAACAACAGAGGAATATGTTGATGACTTATCCAAAGTGGCT	217
Qy	197	GCATTGCTTGCTCAAGCGGGTTGACAAGTTGATCTTTTGGTACGATGTGAGATTTTCA	256
Db	218	GCATCGGACTAATGAAATCAATTGATAATTTTGAOCTGAGGACAAATGTTAAGTTTCAA	277
Qy	257	OCTATCGGGTGCCAATGATCATCGGAGAAATCAAGCTTTTGGCGGATGAAGGTACGG	316
Db	278	CATATGCTGTACCAATGATCATCGGAGAAATCGGAGATATTTGGCGGATACAAACCGA	337
Qy	317	TTAAGGTGAGTGGATGTTTAAAGAAACAGCGAATAGGTGGGGGATCAAGGATGAAT	376
Db	338	TCCGGGCTGACGGGTCACTGAGGATATGGGTACAAAGGGCTGCAAGTGAGAGAACGGC	397
Qy	377	TGTACAAGCAATTTGGGGGTTGGGGGACGATCGCAGAGGTGGCAGAGCAGTGGGAATCA	436
Db	398	TGATCAGTGAGACAAGGAGGGAGGGGACTGCTCAGGAGATGGCTAAAGAGCTTGAAGTGT	457
Qy	437	CGCGGAGGAGGTAGTCTTTGGCGAAGGCAAGCAGAGGGGGCTTCTCCATCGATGAGA	496
Db	458	CCCATGAAGAAATGGTTTTGGGGTTGAGGGCAATTCAGATGCTGTATCTTGTTTGAGC	517
Qy	497	CGGTTTTTGAAGATGAGGGGATCCCATCACACTGATGATCAGATAGCGGATGA	551

Db 518 CGATTTCACATGACCGAGGATCCGATTATGTTCATGGATCAAATCAGCGATGA 572

RESULT 7

ADH97013

ID ADH97013 standard; DNA; 1110 BP.

XX

AC ADH97013;

XX

DT 06-MAY-2004 (first entry)

XX

DE S. pneumoniae RNA polymerase sigma-70 factor gene #2.

XX

KW anti bacterial; anti inflammatory; gastrointestinal; anti ulcer;  
KW anti diarrhoeic; ophthalmological; enzyme inhibitor; antisense therapy;  
KW vaccine; microbial target; modulator; furuncle; pneumonia; gastritis;  
KW peptic ulcer disease; diarrhoea; meningitis; bacteraemia; conjunctivitis;  
KW toxic shock syndrome; gene; ds.

XX

CS Streptococcus pneumoniae.

XX

PN WC2003102190-A2.

XX

PD 11-DEC-2003.

XX

PF 02-JUN-2003; 2003WC000786.

XX

PR 31-MAY-2002; 2002US-0384634P.

PR 31-MAY-2002; 2002US-0385157P.

PR 04-JUN-2002; 2002US-0385542P.

PR 04-JUN-2002; 2002US-0385611P.

PR 04-JUN-2002; 2002US-0385747P.

PR 04-JUN-2002; 2002US-0385750P.

PR 04-JUN-2002; 2002US-0385752P.

PR 04-JUN-2002; 2002US-0385773P.

PR 04-JUN-2002; 2002US-0385780P.

PR 04-JUN-2002; 2002US-0385785P.

PR 04-JUN-2002; 2002US-0385797P.

PR 05-JUN-2002; 2002US-0385962P.

PR 05-JUN-2002; 2002US-0386022P.

PR 05-JUN-2002; 2002US-0386024P.

PR 05-JUN-2002; 2002US-0386087P.

PR 05-JUN-2002; 2002US-0386141P.

PR 05-JUN-2002; 2002US-0386350P.

PR 05-JUN-2002; 2002US-0386586P.

PR 06-JUN-2002; 2002US-0386368P.

PR 06-JUN-2002; 2002US-0386369P.

PR 06-JUN-2002; 2002US-0386436P.

PR 06-JUN-2002; 2002US-0386441P.

PR 06-JUN-2002; 2002US-0386528P.

PR 06-JUN-2002; 2002US-0386573P.

PR 06-JUN-2002; 2002US-0386834P.

PR 31-JUL-2002; 2002US-0399839P.

PR 31-JUL-2002; 2002US-0399861P.

PR 31-JUL-2002; 2002US-0399869P.

PR 31-JUL-2002; 2002US-0399707P.

PR 31-JUL-2002; 2002US-0399883P.

PR 31-JUL-2002; 2002US-0399884P.

PR 31-JUL-2002; 2002US-0399885P.

PR 01-AUG-2002; 2002US-0400154P.

PR 01-AUG-2002; 2002US-0400230P.

PR 01-AUG-2002; 2002US-0400268P.

PR 01-AUG-2002; 2002US-0400363P.

PR 01-AUG-2002; 2002US-0400365P.

PR 01-AUG-2002; 2002US-0400374P.

PR 01-AUG-2002; 2002US-0400380P.

PR 01-AUG-2002; 2002US-0400433P.

PR 01-AUG-2002; 2002US-0400434P.

PR 01-AUG-2002; 2002US-0400436P.

PR 01- AUG- 2002; 2002US- 0400442P.  
PR 01- AUG- 2002; 2002US- 0400463P.

XX  
PA (AFFI - ) AFFI NI UM PHARM INC.  
XX

PI Edwards A, Dharansi A, Vedadi M, Vallee F, Awrey D, Beattie B;  
PI Richards D, Domagal a M, Mansoury K, Virag C, Buzadzija K;  
PI Mcdonal d M, Houston S, Arrowsmith C, Ouyang H, Nethery K, Ng I;  
PI Kanagarajah D;  
XX

DR WPI : 2004- 071165/ 07.  
DR P- PSDB; ADH97014.  
XX

PT Compositions comprising recombinant polypeptide targets for pathogenic  
PT bacteria, useful for designing modulators for preventing or treating a  
PT disease or disorder associated with the species of origin for the  
PT polypeptide.  
XX

PS Claim 23; SEQ ID NO 204; 606pp; English.  
XX

CC The invention relates to novel compositions (I) comprising isolated,  
CC recombinant polypeptides, amino acid sequences having at least about 95%  
CC identity with these or an amino acid sequence encoded by a polynucleotide  
CC that hybridizes under stringent conditions to the complementary strand of  
CC the polynucleotide encoding these polypeptides. The compositions and  
CC polypeptides are useful as microbial targets for designing modulators for  
CC the prevention or treatment of a disease or disorder associated with the  
CC species of origin for the polypeptide, e.g. furuncle, pneumonia,  
CC gastritis, peptic ulcer disease, diarrhoea, meningitis, bacteraemia,  
CC conjunctivitis or toxic shock syndrome. The polypeptides are also useful  
CC for diagnosing a patient suffering from a disease or disorder of a  
CC pathogenic species, or for monitoring the effectiveness of an anti-  
CC pathogenic treatment. This sequence corresponds to one of the DNA  
CC sequences of the invention  
XX

SQ Sequence 1110 BP; 341 A; 211 C; 267 G; 291 T; 0 U; 0 Q her;

Query Match 10.3% Score 78; DB 12; Length 1110;  
Best Local Similarity 50.7% Pred. No. 8.1e-14;  
Matches 214; Conservative 0; Mismatches 205; Indels 3; Gaps 1;

Qy 31 CCATTTCTGACCAATGACCAAGTGAAAGATTTGATAGCCAAGAGCCAGCTGGCGATACG 90  
Db 328 CCTCTCTTGACCAATGACGAGGAGAAAGAGTTGCACTGCCTGTTGAAGCTGGTGATATC 387  
Qy 91 GATGCAAGCTGAGCTTCTCGTGAATAGCAATATCAGACTGGTCTGGTGGTGGTGGTGGTGGT 150  
Db 388 GAGGCAAAACAAAGCTCTTGCGGAAAGCAATCTTGTTTGGTGGTGGTGGTGGTGGTGGTGGT 447  
Qy 151 TTTATCAAGCGGGGATGAGCGGATGATTTGTTTGCATGCGTGGTGGTGGTGGTGGTGGTGGT 210  
Db 448 TATGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGT 507  
Qy 211 AAGGCGGTTGACAAAGTTCGATCTTTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGT 270  
Db 508 AAGGCGGTTGACAAAGTTCGATCTTTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGT 567  
Qy 271 ATGATCATCGAGAAAT---TCAAGCGCTTTTGGGCGATGAGCGTACGGTAAAGGTCAGT 327  
Db 568 TGATTTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 627  
Qy 328 CGATGTTAAAGAAACAGGGAATAGGTTGGGCGATCAAGGATGAATTTGACAAGCAA 387  
Db 628 GTTCACATGGTGAAGCTATCAATAAATGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 687  
Qy 388 TTTGGGCGGTTGGGCGGATGCGGAGGTTGGGCGGATGCGGAGGTTGGGCGGATGCGGAGGTT 447  
Db 688 TTTGGGCGGATGCGGAGGTTGGGCGGATGCGGAGGTTGGGCGGATGCGGAGGTTGGGCGGAT 747

Qy 448 GT 449  
 Db 748 GT 749

## RESULT 8

AEC13368

ID AEC13368 standard; DNA; 1110 BP.

XX

AC AEC13368;

XX

DT 20- OCT- 2005 (first entry)

XX

DE Streptococcus pneumoniae RNA polymerase sigma-70 factor gene.

XX

KW protein purification; antibacterial; antimicrobial; infection;

KW

drug screening; RNA polymerase sigma-70 factor; gene; ss.

XX

CS Streptococcus pneumoniae.

XX

PN US2005181464- A1.

XX

PD 18- AUG- 2005.

XX

PF 29- SEP- 2004; 2004US- 00953901.

XX

PR 04- APR- 2002; 2002US- 0369819P.

PR

PR 04- APR- 2002; 2002US- 0369826P.

PR

PR 04- APR- 2002; 2002US- 0369831P.

PR

PR 04- APR- 2002; 2002US- 0370060P.

PR

PR 08- APR- 2002; 2002US- 0370681P.

PR

PR 08- APR- 2002; 2002US- 0370806P.

PR

PR 08- APR- 2002; 2002US- 0370852P.

PR

PR 08- APR- 2002; 2002US- 0370868P.

PR

PR 09- APR- 2002; 2002US- 0370959P.

PR

PR 09- APR- 2002; 2002US- 0370978P.

PR

PR 09- APR- 2002; 2002US- 0371008P.

PR

PR 09- APR- 2002; 2002US- 0371009P.

PR

PR 09- APR- 2002; 2002US- 0371014P.

PR

PR 09- APR- 2002; 2002US- 0371025P.

PR

PR 09- APR- 2002; 2002US- 0371064P.

PR

PR 09- APR- 2002; 2002US- 0371065P.

PR

PR 09- APR- 2002; 2002US- 0371094P.

PR

PR 09- APR- 2002; 2002US- 0371114P.

PR

PR 09- APR- 2002; 2002US- 0371180P.

PR

PR 09- APR- 2002; 2002US- 0371189P.

PR

PR 31- MAY- 2002; 2002US- 0384634P.

PR

PR 31- MAY- 2002; 2002US- 0385157P.

PR

PR 04- JUN- 2002; 2002US- 0385542P.

PR

PR 04- JUN- 2002; 2002US- 0385611P.

PR

PR 04- JUN- 2002; 2002US- 0385747P.

PR

PR 04- JUN- 2002; 2002US- 0385750P.

PR

PR 04- JUN- 2002; 2002US- 0385752P.

PR

PR 04- JUN- 2002; 2002US- 0385773P.

PR

PR 04- JUN- 2002; 2002US- 0385780P.

PR

PR 04- JUN- 2002; 2002US- 0385785P.

PR

PR 04- JUN- 2002; 2002US- 0385797P.

PR

PR 05- JUN- 2002; 2002US- 0385962P.

PR

PR 05- JUN- 2002; 2002US- 0386022P.

PR

PR 05- JUN- 2002; 2002US- 0386024P.

PR

PR 05- JUN- 2002; 2002US- 0386087P.

PR

PR 05- JUN- 2002; 2002US- 0386141P.

PR

PR 05- JUN- 2002; 2002US- 0386350P.

PR

PR 05- JUN- 2002; 2002US- 0386586P.

PR

PR 06- JUN- 2002; 2002US- 0386368P.

PR

PR 06- JUN- 2002; 2002US- 0386369P.

PR

PR 06- JUN- 2002; 2002US- 0386436P.

PR

PR 06- JUN- 2002; 2002US- 0386441P.

PR

PR 06- JUN- 2002; 2002US- 0386528P.

PR 06-JUN-2002; 2002US 0386573P.  
 PR 06-JUN-2002; 2002US 0386834P.  
 PR 31-JUL-2002; 2002US 0399839P.  
 PR 31-JUL-2002; 2002US 0399861P.  
 PR 31-JUL-2002; 2002US 0399969P.  
 PR 31-JUL-2002; 2002US 0399970P.  
 PR 31-JUL-2002; 2002US 0399983P.  
 PR 31-JUL-2002; 2002US 0399984P.  
 PR 31-JUL-2002; 2002US 0399985P.  
 PR 01-AUG-2002; 2002US 0400154P.  
 PR 01-AUG-2002; 2002US 0400230P.  
 PR 01-AUG-2002; 2002US 0400268P.  
 PR 01-AUG-2002; 2002US 0400363P.  
 PR 01-AUG-2002; 2002US 0400365P.  
 PR 01-AUG-2002; 2002US 0400374P.  
 PR 01-AUG-2002; 2002US 0400380P.  
 PR 01-AUG-2002; 2002US 0400433P.  
 PR 01-AUG-2002; 2002US 0400434P.  
 PR 01-AUG-2002; 2002US 0400436P.  
 PR 01-AUG-2002; 2002US 0400442P.  
 PR 01-AUG-2002; 2002US 0400463P.  
 PR 04-APR-2003; 2003WO CA000465.  
 PR 08-APR-2003; 2003WO CA000482.  
 PR 08-APR-2003; 2003WO CA000483.  
 PR 02-JUN-2003; 2003WO CA000786.

XX

PA

(AFFI-) AFFINUM PHARM INC.

XX

PI

Edwards A, Dharamsi A, Vedadi M, Alam MZ, Arrowsmith G, Awrey DE;

PI

Beattie B, Buzadzija K, Clarke T, Donagala M, Houston S;

PI

Kanagarajah D, Li Q, Mansoury K, McDonald M, Nethery-Brox K, Ng I;

PI

Qiyang H, Richards D, Vallée F, Virag C;

XX

DR

WPI: 2005-628190/64.

DR

P-PSDB; AEC13369.

XX

PT

Novel crystallized, recombinant bacterial polypeptide, useful as targets for pathogenic bacteria such as *Helicobacter pylori*, *Staphylococcus aureus*, for detecting pathogenic species in biological sample, and in drug design.

PT

XX

PS

Claim 85; SEQ ID NO 204; 637pp; English.

XX

OC

The invention relates to a composition (I) comprising purified polypeptides from bacteria. Also described: (1) a crystallized, recombinant polypeptide comprising an amino acid sequence of (I), where the polypeptide is in crystal form (2) a crystallized complex comprising the crystallized, recombinant polypeptide and a co-factor or a small organic molecule, where the complex is in crystal form and (3) a host cell comprising a nucleic acid encoding a polypeptide of (I), where a culture of the host cell produces at least about 1 mg of the polypeptide per liter of culture and the polypeptide is at least about one-third soluble as measured by gel electrophoresis. (I) can be used as a target for pathogenic bacteria, useful for detecting the presence of a pathogenic species in a biological sample. (I) is useful for monitoring the effectiveness of anti-pathogenic treatments in an individual suffering from a disease or disorder caused by a pathogenic bacteria, such as infections. (I) is also useful in drug design and screening, for identifying inhibitors of (I), for designing a potential compound that is useful for treating or preventing pathogenic diseases or disorders, for assessing the activity of small molecules and other modulators in in vitro assay, and for developing antimicrobial agents. The present sequence represents a *Streptococcus pneumoniae* RNA polymerase sigma-70 factor gene, which is used in an example from the present invention.

XX

SQ

Sequence 1110 BP; 341 A; 211 C; 267 G; 291 T; 0 U; 0 Q other;

Query Match 10.3% Score 78; DB 18; Length 1110;  
 Best Local Similarity 50.7% Fred. No. 8.1e-14;

Matches 214; Conservative 0; Mismatches 205; Indels 3; Gaps 1;

```

Qy      31 CCATTCTGACCAATGACCAAGTGAAAGATTTGATAGCCAAGGCCAAGCTGGCGATACG 90
Db      328 CCTCTCTTGACCAATGAAGAGGAGAAAGAGTTGGCACTGGCTGTTGAAGCTGGTGATATC 387
Qy      91  GATGCACGTGAGCTTCTCGTGAATAGCAATATCAGACTGGTCTGGTCGGTCGACAGGC 150
Db      388  GAGGCCAAACAACGCTCTTGGGAAACGCAATCTTGGTTGGTTGGTTCCATTGGCAACGC 447
Qy      151 TTTATCAACCGGGGATGAGGGGATGATTTGTTTGCATCGGTTGCATTGGCTTGTCT 210
Db      448  TATGTGCGTGGTGGCATGCAAGTCTCTGACTTGATTCAAGAAAGAAATATGGGCTTGATG 507
Qy      211 AAGGCGGTGACCAAGTTGATCTTTGTCAGATGTGAGATTTTGCACCTATGGGGTGGCA 270
Db      508  AAGGCGGTGACCAAGTTGACTATTCTAAGGGTTCAGTTTCACTTATGCAACTTGG 567
Qy      271 ATGATCATCGGAGAAAT- - -TCAACGCTTTTGGCGATGACGGTACGGTTAAGGTCAGT 327
Db      568  TGGATTGTCAGGCTATCACTGTCATTGGGACCAAGCTCGTACCATCGGTATCCCA 627
Qy      328 CGATCGTTAAAGAAACAGCGAATAAGGTGGGGGATCAAAGATGAATTGTACAAGCAA 387
Db      628  GTTCACATGGTTGAAACTATCAATAAATGGTTGTCGACAGCGGAATCTCCTTCAAGAA 687
Qy      388 TTGGGGCGTGGGGGACGATGCGAAGTGGCAGAAGCAGTGGGAATCAGCGGGAGGAA 447
Db      688  TTGGGCAAGATCGACACCAAGACAGATTCCTGAACGAATGATATGACACCTGATAAG 747
Qy      448 GT 449
Db      748 GT 749

```

## RESULT 9

ADK44581

ID ADK44581 standard; DNA; 915 BP.

XX

AC

XX

DT

XX

DE

XX

KW

XX

OS

XX

PN

XX

PD

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PF

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PR

PR

PR

XX

PA

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PI

XX

DR

DR

XX

PT

PT

Streptococcus pneumoniae gene, Seq ID No 1096.

ds; gene; Antibacterial; Gene therapy; Vaccine; Streptococcus pneumoniae.

Streptococcus pneumoniae.

US699703-B1.

02-MAR-2004.

26-MAY-2000; 2000US-00583110.

02-JUL-1997; 97US-0051553P.

12-MAY-1998; 98US-0085131P.

30-JUN-1998; 98US-00107433.

(GENO-) GENOME THERAPEUTICS CORP.

Doucette-Stamm L, Bush D, Zeng Q, Opperman T, Houseweart CE;

WPI: 2004-212399/20.

P-PSDB; ADK47242.

New nucleic acid molecules and polypeptides useful for diagnosing, preventing and treating pathological conditions resulting from bacterial infection, e.g. Streptococcus pneumoniae infection, and in drug



PT screening.  
 XX  
 PS Disclosure; SEQ ID NO 1096; 301pp; English.  
 XX  
 CC The invention relates to isolated *Streptococcus pneumoniae* nucleic acids  
 CC and polypeptides. The nucleic acids and proteins are useful for  
 CC diagnosing, preventing and treating pathological conditions resulting  
 CC from bacterial infection, such as *S. pneumoniae* infection. These may also  
 CC be used for drug screening procedures. The present sequence represents a  
 CC *Streptococcus pneumoniae* nucleic acid of the invention. Note: The  
 CC sequence data for this patent did not appear in the printed specification  
 CC but was obtained in electronic format directly from USPTO at  
 CC seqdata.uspto.gov/sequence.html.  
 XX  
 SQ Sequence 915 BP; 279 A; 175 C; 220 G; 241 T; 0 U; 0 Other;

Query Match 10.1% Score 76.4; DB 12; Length 915;  
 Best Local Similarity 50.5% Pred. No. 2.4e-13;  
 Matches 213; Conservative 0; Mismatches 206; Indels 3; Gaps 1;

```

Qy      31 CCATTTC TGACCAATGACCAAGTGAAAGATTTGATAGCCAAAGGCCAAGCTGGCGATACG 90
Db      328 CCTCTCTTGACCAATGAAGAGGAGAAAGAGTTGCCACTGGCTGTTGAAGCTGGTGATATC 387

Qy      91 GATGCACGTGAGCTTCTCGTGAATAGCAATATCAGACTGGTCTGGTCCGTCCAGCGC 150
Db      388 GAAGCCAAACAAAGCTCTTGCGGAAGCCAAATCTTGTTGGTTGGTTCCATTGCCAAAGCG 447

Qy      151 TTTATCAACCGCGGGTATGAAGCGGATGATTTGTTTCAGATCGGTTGCATTGGCTTGCTC 210
Db      448 TATGTCCGTGGTGGTATGCAGTTCCTTGACTTGATTCAAGAAAGAAATATGGCGTTGATG 507

Qy      211 AAGCGCGTTGACCAAGTTGATCTTTTGGTACGATGTGAGATTTTGCACCTATGGGGTGGCA 270
Db      508 AAGCGCGTTGACCAAGTTGACTATTCTAAAGGGTTCAAGTTTCAACTATGCAACTTGG 567

Qy      271 ATGATCATCGGAGAAATTCACGCTTTTTGGCGATGAAG--GTACGGTTAAGGTCAGT 327
Db      568 TGGATTGTCAGGCTATCACTGGTGCTATTGCAGAACCAAGCTCGTACCATCGGTATCCCA 627

Qy      328 CGATCGTTAAAAGAAACAGCGAATAGGTTGGCGGATCAAAGGATGAATGTACAAGCAA 387
Db      628 GTTCACATGGTTGAAACTATCAATAAATTGGTTGGTGAACAGCGGAATCTCCTTCAAGAA 687

Qy      388 TTTCCCGCGTGGCCCAACGATGCGAAGTGGCAGAAAGCAGTGGGAATCAAGCGCGAGGAA 447
Db      688 TTTGGGCAAGATCGCAACCAAGACAGATTGCTGAACGAATGGATATGACACCTGATAAG 747

Qy      448 GT 449
Db      748 GT 749
  
```

## RESULT 10

AEJ68509

ID AEJ68509 standard; DNA; 915 BP.

XX

AC AEJ68509;

XX

DT 05-OCT-2006 (first entry)

XX

DE *Streptococcus pneumoniae* strain 14453 protein-encoding DNA, SEQ: 1096.

XX

KW Vaccine; diagnosis; drug discovery; protein production;

KW bacterial infection; *Streptococcus pneumoniae* infection;

KW bacterial meningitis; bacterial pneumonia; bacteremia; otitis media;

KW antibacterial; neuroprotective; antiinflammatory; respiratory-gen.;

KW auditory; gene; ds.

XX

OS Streptococcus pneumoniae; strain 14453.

XX Key Location/Qualifiers

FT CDS 1..915

FT /\*tag= a

FT /product= "Streptococcus pneumoniae protein SEQ ID

FT NO:3757"

XX US7074914-B1.

XX 11-JUL-2006.

XX 30-DEC-2004; 2004US-00028099.

PR 02-JUL-1997; 97US-0051553P.

PR 12-MAY-1998; 98US-0085131P.

PR 30-JUN-1998; 98US-00107433.

PR 26-MAY-2000; 2000US-00583110.

PR 14-AUG-2003; 2003US-00640833.

XX (SNFI) SANOFI PASTEUR LTEE.

XX Doucette-Stamm L, Bush D, Zeng Q, Opperman T, Houseweart CE;

DR WPI; 2006-500481/51.

DR P-PSDB; AEJ71170.

XX New isolated nucleic acid and polypeptide from Streptococcus pneumoniae, useful for diagnosing, preventing, or treating pathological conditions resulting from bacterial infections, e.g. S. pneumoniae infection.

XX Example; SEQ ID NO 1096; 29pp; English.

XX The invention relates to an isolated nucleic acid, especially (AEJ68056), which encodes the Streptococcus pneumoniae protein of AEJ70717. This nucleic acid is one of 2661 disclosed protein-encoding nucleic acids (AEJ67414-AEJ70074) isolated from a Streptococcus pneumoniae strain 14453 genomic library whose predicted products (AEJ70075-AEJ72735) exhibit homology to known prokaryotic, eukaryotic or archaeal open reading frames (ORFs) or proteins. The invention also relates to a recombinant expression vector comprising the nucleic acid of the invention operably linked to a transcription regulatory element; and a host cell comprising the recombinant expression vector. The Streptococcus pneumoniae nucleic acids and proteins of the invention are useful for diagnosing, preventing, or treating pathological conditions resulting from bacterial infections, especially infections caused by Streptococcus pneumoniae such as meningitis, bacteremia, pneumonia and otitis media. They may also be used in vaccine compositions for the treatment of Streptococcus pneumoniae infections and as targets for antibacterial drugs. Additionally the nucleic acids are useful in the production of commercially important proteins such as enzymes for use in fermentation reactions or in the production of commercially useful metabolites. The present sequence represents a Streptococcus pneumoniae strain 14453 protein-encoding nucleic acid which was identified in the exemplification of the invention. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from the US patent office at seqdata.uspto.gov/sequence.html?DocID=7074914B1.

XX Sequence 915 BP; 279 A; 175 C; 220 G; 241 T; 0 U; 0 Q her;

XX Query Match 10.1% Score 76.4; DB 21; Length 915;  
Best Local Similarity 50.5% Pred. No. 2.4e-13;  
Matches 213; Conservative 0; Mismatches 206; Indels 3; Gaps 1;

Qy 31 CCATTTCACCAATGACCAAGTGAAGATTTGATAGCCAGAGCCAGCTGGCGATACG 90  
Db 328 CCTCTCTTACCAATGACGAGGAGAAAGATTGGCACTGCTGTGTAGCTGTGATATC 387

```

Qy      91  GATGCACGTGACCTTCTCGTGAATAGCAATATCAGACTGGTCTGGTGGTGGTCCAGGCG 150
Db      388  GAAGCCAAACAAAGCTCTTGGGAAAGCCAAATCTTGGTTTGGTTGTTTCCATTGGCAAAAGCG 447
Qy      151  TTTATCAACCGGCGGTATGAAGGGATGATTTGTTTCAGATCGTTCGATTGGCTTGCTC 210
Db      448  TATGTCTGGTGGTGGTATGCAGTTCCTTGACTTGATTCAAGAAAGAAATATGGGCTTGATG 507
Qy      211  AAGGCGGTGACAAAGTTCGATCTTTCGTACGATGTGAGATTTTCGACCTATGGGGTGGCA 270
Db      508  AAGGCGGTGACAAAGTTCGATCTTTCGTACGATGTGAGATTTTCGACCTATGGGGTGGCA 567
Qy      271  ATGATCATCGGAGAAATCAAGCTTTTTGGGCGATGAAG - - GTACGGTTAAGGTCAGT 327
Db      568  TGGATTCTGCAGGCTATCACTGGTCTATTGCAGACCAAGCTCGTACCATCGTATCCCA 627
Qy      328  CGATCGTTAAAGAAACAGCGAATAAGGTGGGGGATCAAAGATGAATTGACAAGCAA 387
Db      628  GTTCACATGGTTGAAACTATCAATAAATGGTTGGTGAACAGCGGAATCTCCTTCAAGAA 687
Qy      388  TTGGGGGTTGGGGGCAAGATCGCAGAAAGTGGCAGAAAGCTGGGAATCAAGCGGAGGAA 447
Db      688  TTGGGGCAAGATCGCAGACCAAGACAGATTGCTGAACGAATGGATATGACAAGCTGATAAG 747
Qy      448  GT 449
Db      748  GT 749

```

## RESULT 11

AEJ75484

ID AEJ75484 standard; DNA; 915 BP.

XX

AC

XX

DT

XX

DE

XX

KW

KW

KW

KW

KW

XX

OS

XX

FH

FT

FT

FT

FT

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PN

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PD

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PF

XX

PR

PR

PR

PR

PR

XX

FA

XX

PI

XX

Streptococcus pneumoniae strain 14453 protein-encoding DNA, SEQ: 1096.

Vaccine; diagnosis; drug discovery; protein production;  
 bacterial infection; Streptococcus pneumoniae infection;  
 bacterial meningitis; bacterial pneumonia; bacteremia; otitis media;  
 antibacterial; neuroprotective; antiinflammatory; respiratory-gen.;  
 auditory; gene; ds.

Streptococcus pneumoniae; strain 14453.

Key Location/Qualifiers

CDS 1..915

/tag= a

/product= "Streptococcus pneumoniae protein SEQ ID NO.3757"

US7081530-B1.

25-JUL-2006.

30-DEC-2004; 2004US-00028291.

02-JUL-1997; 97US-0051553P.

12-MAY-1998; 98US-0085131P.

30-JUN-1998; 98US-00107433.

26-MAY-2000; 2000US-00583110.

14-AUG-2003; 2003US-00640833.

(SNFI) SANOFI PASTEUR LTEE.

Doucette-Stamm L, Bush D, Zeng Q, Opperman T, Houseweart CE;

DR WPI: 2006-518920/53.  
 DR P-PSDB; AEJ78145.  
 XX  
 PT New isolated *Streptococcus pneumoniae* nucleic acid, useful as a molecular  
 PT target for detecting, diagnosing, preventing, or treating a pathological  
 PT condition resulting from bacterial infection.  
 XX  
 PS Example; SEQ ID NO 1096; 29pp; English.  
 XX  
 CC The invention relates to an isolated nucleic acid, especially (AEJ75443),  
 CC which encodes the *Streptococcus pneumoniae* protein of AEJ78104. This  
 CC nucleic acid is one of 2661 disclosed protein-encoding nucleic acids  
 CC (AEJ74389-AEJ77049) isolated from a *Streptococcus pneumoniae* strain 14453  
 CC genomic library whose predicted products (AEJ77050-AEJ79710) exhibit  
 CC homology to known prokaryotic, eukaryotic or archaeal open reading frames  
 CC (ORFs) or proteins. The invention also relates to a recombinant  
 CC expression vector comprising the nucleic acid of the invention operably  
 CC linked to a transcription regulatory element; and a host cell comprising  
 CC the recombinant expression vector. The *Streptococcus pneumoniae* nucleic  
 CC acids and proteins of the invention are useful for diagnosing,  
 CC preventing, or treating pathological conditions resulting from bacterial  
 CC infections, especially infections caused by *Streptococcus pneumoniae* such  
 CC as meningitis, bacteremia, pneumonia and otitis media. They may also be  
 CC used in vaccine compositions for the treatment of *Streptococcus*  
 CC pneumoniae infections and as targets for antibacterial drugs.  
 CC Additionally the nucleic acids are useful in the production of  
 CC commercially important proteins such as enzymes for use in fermentation  
 CC reactions or in the production of commercially useful metabolites. The  
 CC present sequence represents a *Streptococcus pneumoniae* strain 14453  
 CC protein-encoding nucleic acid which was identified in the exemplification  
 CC of the invention. Note: The sequence data for this patent did not form  
 CC part of the printed specification, but was obtained in electronic format  
 CC directly from the US patent office at  
 CC seqdata.uspto.gov/sequence.html?DocID=7081530B1.  
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 SQ Sequence 915 BP; 279 A; 175 C; 220 G; 241 T; 0 U; 0 Other;

Query Match 10.1% Score 76.4; DB 21; Length 915;  
 Best Local Similarity 50.5% Pred. No. 2.4e-13;  
 Matches 213; Conservative 0; Mismatches 206; Indels 3; Gaps 1;

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Db      388 GAAGCAAAACAAGTCTTGCGGAGGCAATCTTGTTTGGTGGTGGTGGTGGTGGTGGTGGTGGT 447
Qy     151 TTTATCAAGCGGGATGAAGCGGATGATTTGTTTGGATGGTGGTGGTGGTGGTGGTGGTGGT 210
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Db     508 AAGGGCGTGGCAAGTTCGATCTTTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGT 567
Qy     271 ATGATCATCGAGAAATTCAGCGCTTTTGGGCGATGAAG--GTACGGTAAAGTGCAGT 327
Db     568 TGATTTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGT 627
Qy     328 CGATGTTAAAGAAACAGGAAATAGGTTGGGCGATGAAGGATGAATTTGACAAGCAA 387
Db     628 GTTCACATGGTGAAGTATCAATAAATGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGT 687
Qy     388 TTTGGGCGTGGGCGCGATGGCAAGTGGCAAGGAGTGGCAATCAAGCGGAGGAA 447
Db     688 TTTGGGCGAGATGGCAAGGAGGAGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGT 747
    
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Qy 448 GT 449  
Db 748 GT 749

RESULT 12

AEJ82844

ID AEJ82844 standard; DNA; 915 BP.

XX

AC AEJ82844;

XX

DT 19- OCT- 2006 (first entry)

XX

DE Streptococcus pneumoniae strain 14453 protein-encoding DNA, SEQ 1096.

XX

KW Vaccine; diagnosis; drug discovery; protein production;  
KW bacterial infection; Streptococcus pneumoniae infection;  
KW bacterial meningitis; bacterial pneumonia; bacteremia; otitis media;  
KW antibacterial; neuroprotective; antiinflammatory; respiratory-gen.;  
KW auditory; gene; ds.

XX

CS Streptococcus pneumoniae; strain 14453.

XX

FH Key Location/Qualifiers

FT CDS

FT

FT

FT

FT

XX

PN

XX

PD

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PF

XX

PR

PR

PR

PR

PR

XX

PA

XX

PI

XX

DR

DR

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PT

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CC

1..915  
/\*tag= a  
/product= "Streptococcus pneumoniae protein SEQ ID  
NO 3757"

US7098023- B1.

29- AUG- 2006.

30- DEC- 2004; 2004US- 00027878.

02- JUL- 1997; 97US- 0051553P.

12- MAY- 1998; 98US- 0085131P.

30- JUN- 1998; 98US- 00107433.

26- MAY- 2000; 2000US- 00583110.

14- AUG- 2003; 2003US- 00640833.

(SNFI ) SANOFI PASTEUR LTD.

Doucette- Stamm L, Bush D, Zeng Q, Opperman T, Houseweart CE;

WPI ; 2006- 584390/ 60.

P- PSDB; AEJ85505.

New isolated Streptococcus pneumoniae nucleic acid and polypeptide,  
useful as vaccines and as targets for diagnosing, preventing, or treating  
pathological conditions resulting from S. pneumoniae bacterial infection.

Example; SEQ ID NO 1096; 29pp; English.

The invention relates to an isolated nucleic acid, especially (AEJ84330),  
which encodes the Streptococcus pneumoniae protein of AEJ86991. This  
nucleic acid is one of 2661 disclosed protein-encoding nucleic acids  
(AEJ81749-AEJ84409) isolated from a Streptococcus pneumoniae strain 14453  
genomic library whose predicted products (AEJ84410-AEJ87070) exhibit  
homology to known prokaryotic, eukaryotic or archaeal open reading frames  
(ORFs) or proteins. The invention also relates to a recombinant  
expression vector comprising the nucleic acid of the invention operably  
linked to a transcription regulatory element; and a host cell comprising  
the recombinant expression vector. The Streptococcus pneumoniae nucleic  
acids and proteins of the invention are useful for diagnosing,  
preventing, or treating pathological conditions resulting from bacterial  
infections, especially infections caused by Streptococcus pneumoniae such  
as meningitis, bacteremia, pneumonia and otitis media. They may also be  
used in vaccine compositions for the treatment of Streptococcus

CC pneumoniae infections and as targets for antibacterial drugs.  
 CC Additionally the nucleic acids are useful in the production of  
 CC commercially important proteins such as enzymes for use in fermentation  
 CC reactions or in the production of commercially useful metabolites. The  
 CC present sequence represents a Streptococcus pneumoniae strain 14453  
 CC protein-encoding nucleic acid which was identified in the exemplification  
 CC of the invention. Note: The sequence data for this patent did not form  
 CC part of the printed specification, but was obtained in electronic format  
 CC directly from the US patent office at  
 CC seqdata.uspto.gov/sequence.html?DocID=7098023B1.  
 XX  
 SQ Sequence 915 BP; 279 A; 175 C; 220 G; 241 T; 0 U; 0 Other;

Query Match	10.1%	Score 76.4;	DB 21;	Length 915;
Best Local Similarity	50.5%	Pred. No. 2.4e-13;		
Matches 213;	Conservative 0;	Mismatches 206;	Indels 3;	Gaps 1;

  

Qy	31	CCATTTCTGACCAATGACCAAGTGAAGATTTGATAGCCAAGGCCAAGCTGGCGATACG	90
Db	328	OCTCTCTTGACCAATGAAGAGGAGAAAGAGTTGGCACTGGCTGTTGAAGCTGGTGATATC	387
Qy	91	GATGCACTGAGCTTCTCGTGAATAGCAATATCAGACTGGTCTGGTGGTGGTCCAGGCG	150
Db	388	GAAGCCAAACAAGCTCTTGGGAAAGCAATCTTGGTTGGTTGGTTCCATTGGCAAAAGCG	447
Qy	151	TTTATCAACCGCGGGTATGAAGCGGATGATTTGTTTGCAGATGGTTCGATTGGCTTGCTC	210
Db	448	TATGTGGTGGTGGTATGCAAGTTCCTTGACTTGATTCAAGAAAGAAATATGGGCTTGATG	507
Qy	211	AAGCGCGTTGACAAGTTGATCTTTGGTACGATGTGAGATTTTGGACCTATGGGGTGGCA	270
Db	508	AAGCGCGTTGACAAGTTGACTATTTCTAAGGGTTCAAGTTTCAACTTATGCAACTTGG	567
Qy	271	ATGATCATCGGAGAAATTCACGCTTTTGGCGGATGAAG--GTACGGTTAAGGTCAGT	327
Db	568	TGGATTGCTGACGGCTATCACTGGTGTCTATTGCAGAACCAAGCTCGTACCATCGGTATCCCA	627
Qy	328	CGATGGTTAAAAGAAACAGCGAATAAGGTGGCGGATCAAGGATGAATTGTACAACCAA	387
Db	628	GTTTCACATGGTTGAACTATCAATAAATGGTTGGTGAACAGCGGAATCTCCTTCAAGAA	687
Qy	388	TTGGCGCGTGGCGGACGATGCGAAGTGGCAGAACGAGTGGGAATCAAGCGGAGGAA	447
Db	688	TTGGGCAAGATGCGACACAGAACAGATTTGGTGAAGCAATGGATATGACACCTGATAAG	747
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## RESULT 13

AEL05163

ID AEL05163 standard; DNA; 915 BP.

XX

AC AEL05163;

XX

DT 30-NOV-2006 (first entry)

XX

DE Streptococcus pneumoniae strain 14453 protein-encoding DNA, SEQ. 1096.

XX

KW Vaccine; diagnosis; drug discovery; protein production;  
 KW bacterial infection; Streptococcus pneumoniae infection;  
 KW bacterial meningitis; bacterial pneumonia; bacteremia; otitis media;  
 KW antibacterial; neuroprotective; antiinflammatory; respiratory-gen.;  
 KW auditory; gene; ds.

XX

CS Streptococcus pneumoniae; strain 14453.

XX

FH Key Location/Qualifiers

FT CDS 1. .915  
 FT /\*tag= a  
 FT /product= "Streptococcus pneumoniae protein SEQ ID  
 FT NO:3757"  
 XX  
 PN US7115731-B1.  
 XX  
 PD 03- OCT-2006.  
 XX  
 PF 30- DEC-2004; 2004US-00027399.  
 XX  
 PR 02- JUL-1997; 97US-0051553P.  
 PR 12- MAY-1998; 98US-0085131P.  
 PR 30- JUN-1998; 98US-00107433.  
 PR 26- MAY-2000; 2000US-00583110.  
 PR 14- AUG-2003; 2003US-00640833.  
 XX  
 PA (SNFI) SANOFI PASTEUR LTD.  
 XX  
 PI Doucette-Stamm L, Bush D, Zeng Q, Opperman T, Houseweart CE;  
 XX  
 DR WPI; 2006-744050/76.  
 DR P-PSDB; AEL07824.  
 XX  
 PT New nucleic acid encoding Streptococcus pneumoniae polypeptide, useful  
 PT for detecting, preventing, and treating pathological conditions resulting  
 PT from bacterial infection.  
 XX  
 PS Example; SEQ ID NO 1096; 29pp; English.  
 XX  
 CC The invention relates to an isolated nucleic acid, especially (AEL05123),  
 CC which encodes the Streptococcus pneumoniae protein of AEL07784. This  
 CC nucleic acid is one of 2661 disclosed protein-encoding nucleic acids  
 CC (AEL04068-AEL06728) isolated from a Streptococcus pneumoniae strain 14453  
 CC genomic library whose predicted products (AEL06729-AEL09389) exhibit  
 CC homology to known prokaryotic, eukaryotic or archaeal open reading frames  
 CC (ORFs) or proteins. The invention also relates to a recombinant  
 CC expression vector comprising the nucleic acid of the invention operably  
 CC linked to a transcription regulatory element; and a host cell comprising  
 CC the recombinant expression vector. The Streptococcus pneumoniae nucleic  
 CC acids and proteins of the invention are useful for diagnosing,  
 CC preventing, or treating pathological conditions resulting from bacterial  
 CC infections, especially infections caused by Streptococcus pneumoniae such  
 CC as meningitis, bacteremia, pneumonia and otitis media. They may also be  
 CC used in vaccine compositions for the treatment of Streptococcus  
 CC pneumoniae infections and as targets for antibacterial drugs.  
 CC Additionally the nucleic acids are useful in the production of  
 CC commercially important proteins such as enzymes for use in fermentation  
 CC reactions or in the production of commercially useful metabolites. The  
 CC present sequence represents a Streptococcus pneumoniae strain 14453  
 CC protein-encoding nucleic acid which was identified in the exemplification  
 CC of the invention. Note: The sequence data for this patent did not form  
 CC part of the printed specification, but was obtained in electronic format  
 CC directly from the US patent office at  
 CC seqdata.uspto.gov/sequence.html?DocID=7115731B1.  
 XX  
 SQ Sequence 915 BP; 279 A; 175 C; 220 G; 241 T; 0 U; 0 Q her;

Query Match 10.1% Score 76.4; DB 21; Length 915;  
 Best Local Similarity 50.5% Pred. No. 2.4e-13;  
 Matches 213; Conservative 0; Msnatches 206; Indels 3; Gaps 1;

Qy 31 CCATTTCTGACCAATGACCAAGTGAAGATTTGATAGCAAGAGCCAGCTGCGATACG 90  
 Db 328 CCTCTCTTGACCAATGAGAGAGAAAGAGTTGGCACTGCGTGTTGAAGCTGTTGATATC 387  
 Qy 91 GATGCAAGTGACCTTCTCGTGAATAGCAATATCAGACTGGTCTGGTGGGCGTGGCCAGCGC 150  
 Db 388 GAAGCCAAACAGCTCTTGGGAGGCAATCTTGGTTGGTGGTTTCCATTGCCAAAGC 447

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 Db 628 GTTCACATGGTTGAACTATCAATAAATTGGTTGCTGAACAGCGGAATCTCTTCAAGAA 687  
 Qy 388 TTGCGCGCTGCGCGCAAGATCGCAGAAAGTGGCAGAAAGCAGTGGGAATCAAGCGGAGAA 447  
 Db 688 TTGGGCAAGATCGACACCAAGAACAGATTGCTGAACGAATGGATATGACAAGTATAAG 747  
 Qy 448 GT 449  
 Db 748 GT 749

## RESULT 14

AEL12413

ID AEL12413 standard; DNA; 915 BP.

XX

AC AEL12413;

XX

DT 28-DEC-2006 (first entry)

XX

DE Streptococcus pneumoniae strain 14453 protein-encoding DNA, SEQ 1096.

XX

KW Vaccine; diagnosis; drug discovery; protein production;  
 KW bacterial infection; Streptococcus pneumoniae infection;  
 KW bacterial meningitis; bacterial pneumonia; bacteremia; otitis media;  
 KW antibacterial; neuroprotective; antiinflammatory; respiratory-gen.;  
 KW auditory; gene; ds.

XX

OS Streptococcus pneumoniae; strain 14453.

XX

FH Key Location/Qualifiers

CDS 1..915

FT

FT /tag= a

FT /product= "Streptococcus pneumoniae protein SEQ ID

FT NO.3757"

XX

PN US7122368-B1.

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PD 17-OCT-2006.

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PF 30-DEC-2004; 2004US-00027877.

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PR 02-JUL-1997; 97US-0051553P.

PR 12-MAY-1998; 98US-0085131P.

PR 30-JUN-1998; 98US-00107433.

PR 26-MAY-2000; 2000US-00583110.

PR 14-AUG-2003; 2003US-00640833.

XX

FA (SNFI) SANOFI PASTEUR LTD.

XX

FI Doucette-Stamm L, Bush D, Zeng Q, Opperman T, Houseweart CE;

XX

DR WPI: 2006-812612/82.

DR P-PSDB; AEL15074.

XX





## RESULT 15

AEL50821

ID AEL50821 standard; DNA; 915 BP.

XX

AC AEL50821;

XX

DT 28-DEC-2006 (first entry)

XX

DE Streptococcus pneumoniae strain 14453 protein-encoding DNA, SEQ 1096.

XX

KW Vaccine; diagnosis; drug discovery; protein production;  
 KW bacterial infection; Streptococcus pneumoniae infection;  
 KW bacterial meningitis; bacterial pneumonia; bacteremia; otitis media;  
 KW antibacterial; neuroprotective; antiinflammatory; respiratory-gen.;  
 KW auditory; gene; ds.

XX

CS Streptococcus pneumoniae; strain 14453.

XX

FH Key Location/Qualifiers

FT CDS

FT

FT

FT

FT

XX

PN US7129340-B1.

XX

PD 31-OCT-2006.

XX

PF 30-DEC-2004; 2004US-00028457.

XX

PR 02-JUL-1997; 97US-0051553P.

PR 12-MAY-1998; 98US-0085131P.

PR 30-JUN-1998; 98US-00107433.

PR 26-MAY-2000; 2000US-00583110.

PR 14-AUG-2003; 2003US-00640833.

XX

PA (SNFI) SANOFI PASTEUR LTD.

XX

PI Doucette-Stamm L, Bush D, Zeng Q, Opperman T, Houseweart CE;

XX

DR WPI; 2006-812716/82.

DR P-PSDB; AEL53482.

XX

PT New isolated nucleic acid and polypeptide isolated from Streptococcus

PT pneumoniae, useful as components of antibacterial vaccines, and for

PT diagnosing or treating S. pneumoniae and other Streptococcus infections.

XX

PS Example; SEQ ID NO 1096; 29pp; English.

XX

OC The invention relates to an isolated nucleic acid, especially (AEL52290),

OC which encodes the Streptococcus pneumoniae protein of AEL54951. This

OC nucleic acid is one of 2661 disclosed protein-encoding nucleic acids

OC (AEL49726-AEL52386) isolated from a Streptococcus pneumoniae strain 14453

OC genomic library whose predicted products (AEL52387-AEL55047) exhibit

OC homology to known prokaryotic, eukaryotic or archaeal open reading frames

OC (ORFs) or proteins. The invention also relates to a recombinant

OC expression vector comprising the nucleic acid of the invention operably

OC linked to a transcription regulatory element; and a host cell comprising

OC the recombinant expression vector. The Streptococcus pneumoniae nucleic

OC acids and proteins of the invention are useful for diagnosing,

OC preventing, or treating pathological conditions resulting from bacterial

OC infections, especially infections caused by Streptococcus pneumoniae such

OC as meningitis, bacteremia, pneumonia and otitis media. They may also be

OC used in vaccine compositions for the treatment of Streptococcus

OC pneumoniae infections and as targets for antibacterial drugs.

OC Additionally the nucleic acids are useful in the production of

OC commercially important proteins such as enzymes for use in fermentation

reactions or in the production of commercially useful metabolites. The present sequence represents a *Streptococcus pneumoniae* strain 14453 protein-encoding nucleic acid which was identified in the exemplification of the invention. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from the US patent office at [seqdata.uspto.gov/sequence.html?DocID=7129340B1](http://seqdata.uspto.gov/sequence.html?DocID=7129340B1).

Sequence 915 BP; 279 A; 175 C; 220 G; 241 T; 0 U; 0 Q her;

Query Match 10.1% Score 76.4; DB 21; Length 915;  
 Best Local Similarity 50.5% Pred. No. 2.4e-13;  
 Matches 213; Conservative 0; Mismatches 206; Indels 3; Gaps 1;

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Qy      91  GATGCAAGTGAAGCTTCTCGTGAATGCAATATCAGACTGGTCTGGTCCGTCGTCACAGCG 150
Db      388  GAAAGCAAAACAAAGCTCTTGGGAAGCAATCTTGGTGGTGGTTCATTGCCAAACGC 447
Qy     151  TTTATCAACCGCGGGTATGAAGCGGATGATTTGTTTCAGATCGGTTGCATTGCGTCTGCTC 210
Db     448  TATGTGGTGGTGGTATGCAGTTCCTTGACTTGATTCAAGAGGAAATATGGGCTTGATG 507
Qy     211  AAGGCGGTTGACAAGTTGATCTTTGTAAGATGTGAGATTTTGAAGCTATGGGGTGGCA 270
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Qy     271  ATGATCATCGGAGAAATTCAACGCTTTTTTGCGGATGAAG -- GTAAGGTTAAGGTGAGT 327
Db     568  TGATTCGTCAGGCTATCACTGGTCTATTGCAGAACCAAGCTGTAACATCCGATATCCCA 627
Qy     328  GATGCTTTAAAAGAAACAGGGAATTAAGGTGGGGGATCAAGGATGAATGTACAAGCAA 387
Db     628  GTTCACATGGTTGAAACTATCAATAAATTGGTTCGTGAACAGCGGAATCTCCTTCAAGAA 687
Qy     388  TTGCGGCGTGGGCGCAAGATGCGAGAGTGCGAGAGCAGTGGGAATCAAGCGGAGGAA 447
Db     688  TTGGGCAAGATCGGCAACAGAACAGATTGCTGAAAGATGATATGACACTGATAAG 747
Qy     448  GT 449
Db     748  GT 749

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